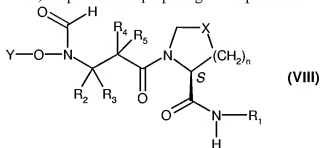


**AMENDMENTS TO THE CLAIMS**

This listing of claims will replace all prior versions, and listings, of claims in the application.

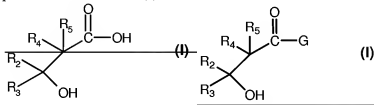
**Listing of Claims:**

Claim 1. (Currently amended) A process for preparing a compound of the formula (VIII)

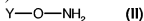


comprising step A:

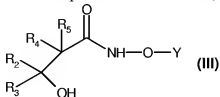
contacting a compound of the formula (I)



with a compound of the formula (II)

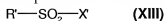


in the presence of a carboxy activating agent, in a suitable solvent  
under conditions suitable to form a compound of the formula (III)

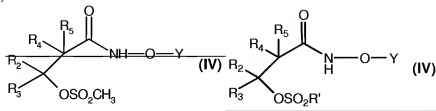


followed by step B:

contacting compound (III) with a compound of the formula (XIII)

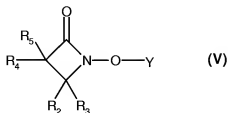


in the presence of a base in a suitable solvent, under conditions suitable to form a compound of the formula (IV)



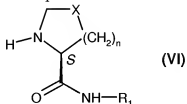
followed by Step C:

contacting compound (IV) with a base in a suitable solvent under conditions suitable to form a compound of the formula (V)

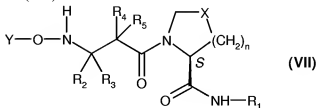


followed by Step D:

contacting compound (V) with a compound of the formula (VI)



in a suitable solvent optionally in the presence of an activator under conditions suitable to form a compound of the formula (VII)



followed by Step E:

contacting compound (VII) with a formylating agent in a suitable solvent under conditions suitable to form compound (VIII);

wherein

Y is a hydroxy protecting group;

Each of  $R_2$ ,  $R_3$ ,  $R_4$  and  $R_5$ , independently, is hydrogen or an aliphatic group, or ( $R_2$  and  $R_3$ ) and/or ( $R_4$  and  $R_5$ ) collectively form a  $C_{4-7}$  cycloalkyl;

X is  $-CH_2-S-$ ,  $CH(OH)-$ ,  $CH(OR)-$ ,  $CH(SH)-$ ,  $CH(SR)-$ ,  $CF_2-$ ,  $C=N(OR)-$  or  $CH(F)-$ ;

wherein

$R$  is alkyl;

G is  $-OH$  or  $-O^{\ominus}M^{\oplus}$ , wherein M is a metal or an ammonium moiety;

$R_1$  is aryl or heteroaryl;

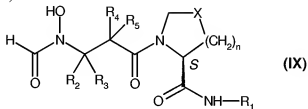
$X'$  is halo;

$R'$  is alkyl or aryl; and

n is 1 0 to 3, provided that when n is 0, X is  $-CH_2-$ .

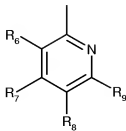
Claim 2. (Original) The process of Claim 1, followed by additional Step F which comprises contacting the compound of formula (VIII), wherein  $R_1$  is heteroaryl having an *N* heteroatom, with an oxidizing agent to form the corresponding *N*-oxide derivative.

Claim 3. (Original) The process of Claim 1, followed by the additional step of removing the hydroxyl-protecting group by contacting compound (VIII) with a palladium catalyst to form the compound of formula (IX)



wherein  $R_1$ ,  $R_2$ ,  $R_3$ ,  $R_4$ ,  $R_5$ , X and n are as defined above.

Claim 4. (Original) The process of Claim 1, wherein each of  $R_2$ ,  $R_3$  and  $R_5$  is hydrogen;  $R_4$  is butyl; X is  $-CH_2-$ ; n is 1; Y is benzyl or *t*-butyldimethylsilyl; and  $R_1$  is of the formula



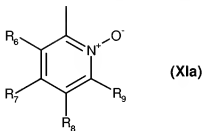
wherein

- R<sub>6</sub> and R<sub>9</sub> are hydrogen;  
R<sub>7</sub> is hydrogen or C<sub>1-7</sub>alkyl; and  
R<sub>8</sub> is hydrogen, halogen or C<sub>1-7</sub>alkyl.

Claim 5. (Currently amended) The process of Claim 4 [[2]], wherein R<sub>7</sub> is hydrogen; and R<sub>8</sub> is fluoro.

Claim 6. (Currently amended) The process of Claim 4 [[2]], wherein R<sub>7</sub> is C<sub>1-7</sub> alkyl; and R<sub>8</sub> is hydrogen.

Claim 7. (Original) The process of Claim 1, wherein R<sub>1</sub> is of the formula (XIa)



wherein

- R<sub>6</sub>, R<sub>7</sub> and R<sub>9</sub> are hydrogen; and  
R<sub>8</sub> is halogen or C<sub>1-7</sub>alkyl.

Claim 8. (Original) The process of Claim 7, wherein R<sub>8</sub> is fluoro.

Claim 9. (Original) The process of Claim 1, carried out at a temperature of about 0°C to about 80°C, a pH of about 2 to about 12, and in one or more solvents selected from the group consisting of dioxane, methylene chloride, dichloromethane, toluene, acetone, methyl ethyl ketone, THF, isopropyl acetate, DMF and an alcohol.

Claims 10 – 37. (Cancelled)